## Amendment to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

claims 1, 2, 5-7, 9, 11 and 27 have been amended; and claims 23 and 24 have been canceled.

## Listing of Claims:

Claim 1 (currently amended): A <u>site-specific drug delivery</u> medical device <u>having a coating</u> consisting essentially of a <u>site-specific delivery device for the controlled release of</u> at least one peroxisome proliferator-activated receptor gamma (PPARγ) agonist and at least one biocompatible polymer.

Claim 2 (currently amended): The <u>site-specific drug delivery</u> medical device according to claim 1 wherein said PPAR<sub>Y</sub> agonist is rosiglitazone.

Claim 3 (canceled)

Claim 4 (canceled)

Claim 5 (currently amended): The <u>site-specific drug delivery</u> medical device according to any of claims 1 or 2 wherein said medical device is a stent.

Claim 6 (currently amended): The <u>site-specific drug delivery</u> medical device according to claim 5 wherein said stent is a vascular stent or biliary stent.

Claim 7 (currently amended): The <u>site-specific drug delivery</u> medical device according to claim 6 wherein said vascular stent is provided with a coating consisting essentially of rosiglitazone and at least one biocompatible polymer.

Claim 8 (canceled)

Claim 9 (currently amended): The <u>site-specific drug delivery</u> medical device according to claim 1 wherein said biocompatible polymer is selected from the group consisting of polyvinyl pyrrolidone, polytetrafluoroethylene, poly-L-lactic acid,

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Reply to Office Action mailed May 4, 2006

polycaprolactone, polyethylene glycol, polystyrene, acrylates, polyesters and mixtures thereof

Claim 10 (canceled)

Claim 11 (currently amended): A <u>vascular stent medical device-consisting</u> essentially of a <u>vascular stent and</u>rosiglitazone; and

a polymer selected from the group consisting of polyvinyl pyrrolidone, polytetrafluoroethylene, poly-L-lactic acid, polycaprolactone, polyethylene glycol, polystyrene, acrylates, polyesters and mixtures thereof.

Claims 12-26 (canceled)

Claim 27 (currently amended): The <u>site-specific drug delivery</u> medical device according to claim 7 wherein said biocompatible polymer is selected from the group consisting of polyvinyl pyrrolidone, polytetrafluoroethylene, poly-L-lactic acid, polycaprolactone, polyethylene glycol, polystyrene, acrylates, polyesters and mixtures thereof.